

Reprinted from THE JOURNAL OF CHEMICAL PHYSICS, Vol. 25, No. 5, 1087, November, 1956
Printed in U. S. A.

Multiple-Path Technique for the Determination of Physico-Chemical Data Behind Shock Fronts*

W. HOOKER, M. LAPP, D. WEBER, AND S. S. PENNER

Daniel and Florence Guggenheim Jet Propulsion Center, California Institute of Technology Pasadena, California

(Received August 27, 1956)

IN connection with current experimental studies on carbon formation and gas emissivities at elevated temperatures behind shock fronts, we are utilizing single-path and double-path emission and absorption measurements.¹ For observations behind shock fronts (observation times of less than about 200 μ sec), we have adopted a procedure involving simultaneous spectral recording of an unchopped single-path beam and of a chopped double-path beam.

A schematic diagram of the experimental arrangement for emission studies is sketched in Fig. 1. A slotted air turbine wheel making 120 000 rpm permits frequency modulation of the double-path beam at a rate of 5 μ sec, which is sufficiently rapid for our particular shock-tube applications. The mechanical chopper is to be preferred to a Kerr cell shutter² because the losses are minimized and because a much wider wavelength range can be covered. The single and double-path beams are observed simultaneously on the same detector and are used only to establish the intensity ratio at various wavelengths as a function of time. Hence errors associated with any variations in the receiving system, other than a loss of linearity, will not affect the observed results.

Interpretation of the measurements is based on the fact that, under suitably chosen conditions, the spectral intensity ratio for the double- and single-path experiments is a unique function of the product $P_\lambda p L$. Here P_λ is the spectral absorption coefficient, p equals the partial pressure of emitter or absorber, and L is the cell length. For chemical reaction rate studies, P_λ and L are known through independent calibrations and analysis; hence the partial pressure of emitter (or absorber) p is determined by the experimental data. In gas emissivity measurements, p and L are known and the measured intensity ratio determines the spectral absorption coefficient P_λ .

* The work on chemical reactions is supported by the U. S. Air Force, Office for Advanced Studies, under Contract AF 18(603)-2; the gas emissivity measurements are being performed with support from the U. S. Navy, Office of Naval Research, under Contract Nonr-220(03), NR 015 401.

¹ A similar procedure has been employed previously for emission studies on flames. See S. S. Penner and E. K. Björnerud, *J. Chem. Phys.* **23**, 143 (1955).

² F. Harshbarger, *J. Chem. Phys.* **24**, 1261 (1956).

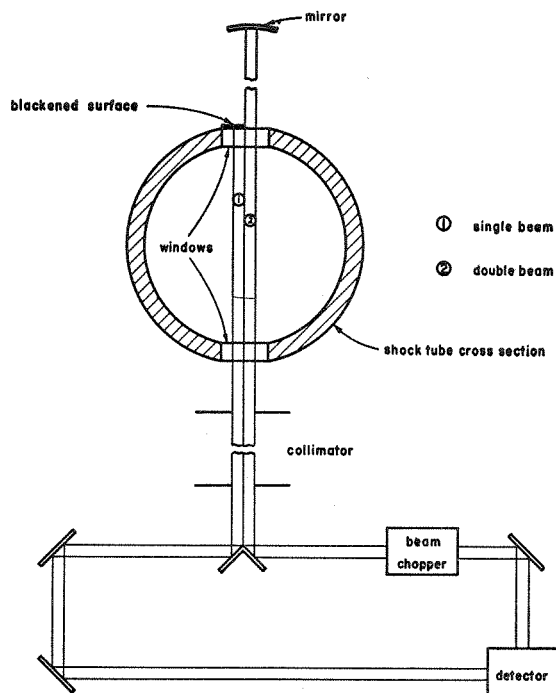


FIG. 1. Schematic diagram of the optical system used for the simultaneous determinations of single-path and double-path intensities. The axis of the shock tube is normal to the plane of the paper.